

(FILE 'HOME' ENTERED AT 22:49:43 ON 09 NOV 2008)

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L1 STRUCTURE UPLOADED
L2 0 S L1 SSS SAM
L3 8 S L1 SSS FULL

FILE 'CAPLUS' ENTERED AT 22:52:48 ON 09 NOV 2008

=> s L3

L4 6 L3

=> d L4 1-6 TI ABS IBIB HITSTR

L4 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN

TI Pharmaceutical compositions based on anticholinergics and additional active ingredients

AB A pharmaceutical compn. comprising an anticholinergic and at least one addnl. active ingredient selected from among corticosteroids, dopamine agonists, PDE-IV inhibitors, NK1-antagonists, endothelin antagonists, antihistamines, and EGFR-kinase inhibitors, processes for prep. them and their use in the treatment of respiratory diseases. Among a no. of compds. prep. was N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-2-[4-[(3-hydroxypropyl)methylamino]piperidin-1-yl]-N-methyl-2-phenylacetamide. Inhalable powders include a formulation contg. tiotropium bromide, budesonide, and lactose.

ACCESSION NUMBER: 2005:586215 CAPLUS

DOCUMENT NUMBER: 143:120526

TITLE: Pharmaceutical compositions based on anticholinergics and additional active ingredients

INVENTOR(S): Pairet, Michel; Pieper, Michael P.; Meade, Christopher
John Montague; Reichl, Richard; Schmelzer, Christel;
Jung, Birgit

PATENT ASSIGNEE(S): Boehringer Ingelheim Pharma GmbH & Co. Kg, Germany

SOURCE: U.S. Pat. Appl. Publ., 50 pp., Cont.-in-part of U.S.
Ser. No. 824,391.

CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

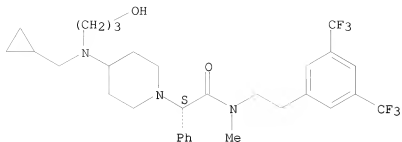
FAMILY ACC. NUM. COUNT: 19

PATENT INFORMATION:

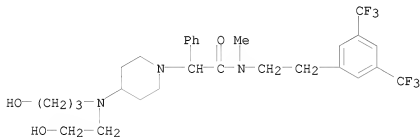
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 20050148562	A1	20050707	US 2004-6940	20041208
DE 10062712	A1	20020620	DE 2000-10062712	20001215
DE 10063957	A1	20020627	DE 2000-10063957	20001220
DE 10110772	A1	20020912	DE 2001-10110772	20010307
DE 10111058	A1	20020912	DE 2001-10111058	20010308
DE 10113366	A1	20020926	DE 2001-10113366	20010320
DE 10138272	A1	20030227	DE 2001-10138272	20010810
US 20020151541	A1	20021017	US 2001-7182	20011019
US 20020183292	A1	20021205	US 2001-86145	20011019
CA 2614631	A1	20020510	CA 2001-2614631	20011023
US 20020137764	A1	20020926	US 2001-40196	20011025
US 20020122773	A1	20020905	US 2001-27662	20011220
DE 10206505	A1	20030828	DE 2002-10206505	20020216
US 20020169181	A1	20021114	US 2002-92116	20020306
US 6620438	B2	20030916		
US 20020193393	A1	20021219	US 2002-93240	20020307

US 20020183347	A1	20021205	US 2002-100659	20020318
US 6608054	B2	20030819		
US 20030158196	A1	20030821	US 2003-360064	20030207
US 20030181478	A1	20030925	US 2003-395777	20030324
US 6890517	B2	20050510		
US 20030203925	A1	20031030	US 2003-413065	20030414
US 20030212075	A1	20031113	US 2003-419358	20030421
US 6696042	B2	20040224		
US 20040024007	A1	20040205	US 2003-613783	20030703
US 20040151770	A1	20040805	US 2004-763894	20040123
US 20040161386	A1	20040819	US 2004-775901	20040210
US 20040176338	A1	20040909	US 2004-776757	20040211
US 20040192675	A1	20040930	US 2004-824391	20040414
US 20050147564	A1	20050707	US 2005-68134	20050228
AU 2008202554	A1	20080703	AU 2008-202554	20080610
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			DE 2000-10054042	A 20001031
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			US 2000-257220P	P 20001221
			US 2000-257221P	P 20001221
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			DE 2001-10111058	A 20010308
			DE 2001-10113366	A 20010320
			US 2001-281653P	P 20010405
			US 2001-281857P	P 20010405
			US 2001-281874P	P 20010405
			DE 2001-10138272	A 20010810
			US 2001-314599P	P 20010824
			US 2001-7182	B1 20011019
			US 2001-86145	B1 20011019
			US 2001-27662	B1 20011220
			DE 2002-10206505	A 20020216
			US 2002-92116	A1 20020306
			US 2002-93240	B1 20020307
			US 2002-100659	A1 20020318
			US 2002-369213P	P 20020401
			US 2003-360064	A2 20030207
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			US 2004-776757	A2 20040211
			US 2004-824391	A2 20040414
			CA 2001-2436540	A3 20011023
			US 2001-40196	B1 20011025
			US 2003-395777	A1 20030324
			AU 2006-202723	A3 20060626
OTHER SOURCE(S): MARPAT 143:120526				
IT	415917-07-6P 457910-79-1P 502422-75-5P			
	RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)			
	(pharmaceutical compns. based on anticholinergics and addnl. active ingredients)			
RN	415917-07-6 CAPLUS			
CN	1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(cyclopropylmethyl)(3-hydroxypropyl)amino]-N-methyl-.alpha.-phenyl-, (.alpha.S)- (CA INDEX NAME)			

Absolute stereochemistry.



RN 457910-79-1 CAPLUS
 CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(2-hydroxyethyl)(3-hydroxypropyl)amino]-N-methyl-.alpha.-phenyl- (CA INDEX NAME)

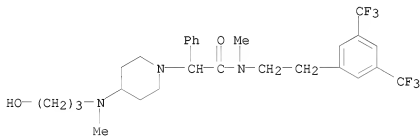


RN 502422-75-5 CAPLUS
 CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(3-hydroxypropyl)methylamino]-N-methyl-.alpha.-phenyl-, (2E)-2-butenedioate (2:3) (salt) (9CI) (CA INDEX NAME)

CM 1

CRN 415916-92-6

CMF C28 H35 F6 N3 O2



CM 2

CRN 110-17-8

CMF C4 H4 O4

Double bond geometry as shown.



L4 ANSWER 2 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN
 TI Pharmaceutical compositions comprising novel anticholinergic agents and
 NK1-receptor antagonists for the treatment of respiratory tract diseases
 AB The invention relates to novel pharmaceutical compns. comprising novel
 anticholinergic agents and NK1-receptor antagonists, method for prodn. and
 use thereof in the treatment of respiratory diseases. Thus an inhalation
 capsule contained (microgram/capsule): 2,2-Diphenylpropionic acid scopoline
 ester methobromide 200; N-[2-(3,5-Bis-trifluoromethylphenyl)-ethyl]-2-(4-
 [(3-hydroxypropyl)methylamino]piperidin-1-yl)-N-methyl-2-phenylacetamide
 150; lactose 12150.

ACCESSION NUMBER: 2004:41273 CAPLUS
 DOCUMENT NUMBER: 140:99643
 TITLE: Pharmaceutical compositions comprising novel
 anticholinergic agents and NK1-receptor antagonists
 for the treatment of respiratory tract diseases
 INVENTOR(S): Pairet, Michel; Meade, Christopher John Montague;
 Pieper, Michael P.
 PATENT ASSIGNEE(S): Boehringer Ingelheim Pharma G.m.b.H. & Co. K.-G.,
 Germany
 SOURCE: PCT Int. Appl., 42 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

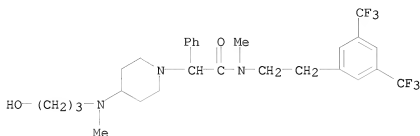
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004004724	A1	20040115	WO 2003-EP6667	20030625
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RR: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
DE 10230750	A1	20040122	DE 2002-10230750	20020709
CA 2491451	A1	20040115	CA 2003-2491451	20030625
AU 2003242754	A1	20040123	AU 2003-242754	20030625
EP 1521580	A1	20050413	EP 2003-762508	20030625
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
JP 2005532378	T	20051027	JP 2004-518565	20030625
US 20040048886	A1	20040311	US 2003-614362	20030707
PRIORITY APPLN. INFO.:			DE 2002-10230750	A 20020709
			US 2002-407758P	P 20020903
			WO 2003-EP6667	W 20030625

OTHER SOURCE(S): MARPAT 140:99643
 IT 415916-92-6
 RL: PEP (Physical, engineering or chemical process); PYP (Physical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
 (pharmaceutical compns. comprising anticholinergic agents and

NK1-receptor antagonists for treatment of respiratory tract diseases)

RN 415916-92-6 CAPLUS

CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(3-hydroxypropyl)methylamino]-N-methyl-.alpha.-phenyl- (CA INDEX NAME)

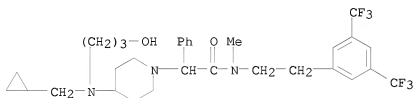


IT 457910-81-5

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(pharmaceutical compns. comprising anticholinergic agents and
NK1-receptor antagonists for treatment of respiratory tract diseases)

RN 457910-81-5 CAPLUS

CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(cyclopropylmethyl)(3-hydroxypropyl)amino]-N-methyl-.alpha.-phenyl- (CA INDEX NAME)

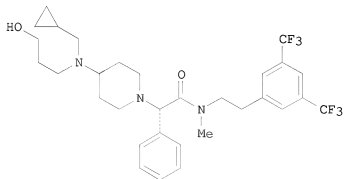


REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 3 OF 6 CAPLUS COPYRIGHT 2008 ACS on SIN

TI Method using NK1 receptor antagonists for the treatment or prevention of atopic dermatitis

GI



AB The invention provides a method for the treatment or prevention of atopic dermatitis, which comprises the administration of an effective amt. of an NK1 receptor antagonist to a patient in need of such treatment, wherein the NK1 receptor antagonist is effective in inhibiting substance P-induced scratching in mice. Compsds. of the invention include e.g. I. Prepn. of selected compds. is described.

ACCESSION NUMBER: 2003:238299 CAPLUS

DOCUMENT NUMBER: 138:248551

TITLE: Method using NK1 receptor antagonists for the

INVENTOR(S): treatment or prevention of atopic dermatitis

PATENT ASSIGNEE(S): Komune, Kunihiro; Ohmura, Tsuyoshi; Satoh, Hisashi

SOURCE: Boehringer Ingelheim International GmbH, Germany

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1295599	A1	20030326	EP 2001-122730	20010921
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
US 20030100565	A1	20030529	US 2002-236824	20020906
WO 2003026658	A1	20030403	WO 2002-EP10502	20020919
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2002333858	A1	20030407	AU 2002-333858	20020919
PRIORITY APPLN. INFO.:			EP 2001-122730	A 20010921
			US 2001-338416P	P 20011115
			WO 2002-EP10502	W 20020919

OTHER SOURCE(S): MARPAT 138:248551

IT 415917-07-6P 415917-12-3P 502422-75-5P

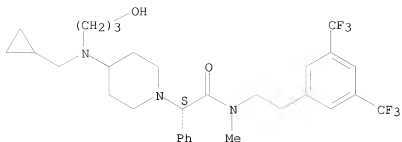
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(NK1 receptor antagonists for treatment or prevention of atopic dermatitis)

RN 415917-07-6 CAPLUS

CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(cyclopropylmethyl)(3-hydroxypropyl)amino]-N-methyl-.alpha.-phenyl-, (.alpha.S)- (CA INDEX NAME)

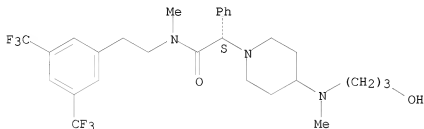
Absolute stereochemistry.



RN 415917-12-3 CAPLUS

CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(3-hydroxypropyl)methylamino]-N-methyl-.alpha.-phenyl-, (.alpha.S)- (CA INDEX NAME)

Absolute stereochemistry.



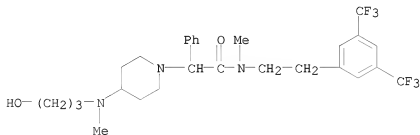
RN 502422-75-5 CAPLUS

CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(3-hydroxypropyl)methylamino]-N-methyl-.alpha.-phenyl-, (2E)-2-butenedioate (2:3) (salt) (9CI) (CA INDEX NAME)

CM 1

CRN 415916-92-6

CMF C28 H35 F6 N3 O2



CM 2

CRN 110-17-8

CMF C4 H4 O4

Double bond geometry as shown.



REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 4 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN
 TI Pharmaceutical compositions based on anticholinergics and NK1-receptor antagonists for the treatment of respiratory tract diseases
 AB The invention discloses pharmaceutical compns. based on anticholinergics and NK1-receptor antagonists, processes for prepn. them, and their use in the treatment of respiratory tract diseases. Prepn. of selected compds. is included.

ACCESSION NUMBER: 2002:869585 CAPLUS

DOCUMENT NUMBER: 137:346202

TITLE: Pharmaceutical compositions based on anticholinergics and NK1-receptor antagonists for the treatment of respiratory tract diseases

INVENTOR(S): Pairet, Michel; Pieper, Michael P.; Meade, Christopher J. M.

PATENT ASSIGNEE(S): Germany

SOURCE: U.S. Pat. Appl. Publ., 18 pp., Cont.-in-part of U. S. Provisional Ser. NO. 281,653.

CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 19

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 20020169181	A1	20021114	US 2002-92116	20020306
US 6620438	B2	20030916		
DE 10111058	A1	20020912	DE 2001-10111058	20010308
US 20030212075	A1	20031113	US 2003-419358	20030421
US 6696042	B2	20040224		
US 20040151770	A1	20040805	US 2004-763894	20040123
US 20050148562	A1	20050707	US 2004-6940	20041208
AU 2008202554	A1	20080703	AU 2008-202554	20080610
PRIORITY APPLN. INFO.:				
			DE 2001-10111058	A 20010308
			US 2001-281653P	P 20010405
			DE 2000-10054042	A 20001031
			US 2000-253613P	P 20001128
			DE 2000-10062712	A 20001215
			DE 2000-10063957	A 20001220
			US 2000-257220P	P 20001221
			US 2000-257221P	P 20001221
			DE 2001-10110772	A 20010307
			DE 2001-10113366	A 20010320
			US 2001-281857P	P 20010405
			US 2001-281874P	P 20010405
			DE 2001-10138272	A 20010810
			US 2001-314599P	P 20010824
			US 2001-7182	B1 20011019
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			US 2001-27662	B1 20011220
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			US 2002-93240	B1 20020307
			US 2002-100659	A1 20020318

US 2002-369213P	P 20020401
US 2003-360064	A2 20030207
US 2003-413065	B2 20030414
US 2003-419358	A1 20030421
US 2003-613783	A2 20030703
US 2004-763894	A2 20040123
US 2004-775901	A2 20040210
US 2004-776757	A2 20040211
US 2004-824391	A2 20040414
AU 2006-202723	A3 20060626

OTHER SOURCE(S): MARPAT 137:346202

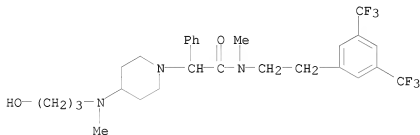
IT 415916-92-6P 415917-07-6P 457910-79-1P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(anticholinergics and NK1-receptor antagonists for treatment of respiratory tract diseases)

RN 415916-92-6 CAPLUS

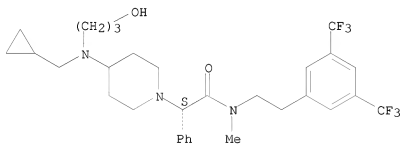
CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(3-hydroxypropyl)methylamino]-N-methyl-.alpha.-phenyl- (CA INDEX NAME)



RN 415917-07-6 CAPLUS

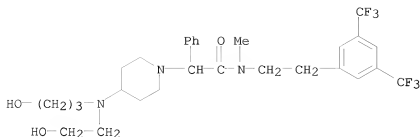
CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(cyclopropylmethyl) (3-hydroxypropyl)amino]-N-methyl-.alpha.-phenyl-, (.alpha.S)- (CA INDEX NAME)

Absolute stereochemistry.

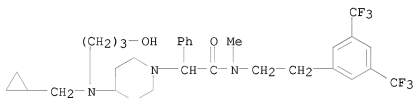


RN 457910-79-1 CAPLUS

CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(2-hydroxyethyl) (3-hydroxypropyl)amino]-N-methyl-.alpha.-phenyl- (CA INDEX NAME)



IT 457910-81-5
 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
 (Biological study); USES (Uses)
 (anticholinergics and NK1-receptor antagonists for treatment of
 respiratory tract diseases)
 RN 457910-81-5 CAPLUS
 CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-
 [(cyclopropylmethyl) (3-hydroxypropyl) amino]-N-methyl-.alpha.-phenyl- (CA
 INDEX NAME)



L4 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN
 TI Inhalant compositions containing anticholinergics and NK1 receptor
 antagonists
 AB The invention relates to drug compns. based on anticholinergics and on NK1
 receptor antagonists, to methods for their prodn., and to their use as
 inhalants for the treatment of respiratory tract diseases. Synthesis of
 NK1 receptor antagonists from the group of
 bis-trifluoromethyl-phenyl-piperidine derivs. are described. The products
 are used in suspension aerosols. Thus a compn. contained (wt./wt.%):
 tiotropium bromide 0.015; NK1 receptor antagonist 0.066; soy lecithin 0.2;
 TG11: TG12 = 2:3 to 100.

ACCESSION NUMBER: 2002:695760 CAPLUS
 DOCUMENT NUMBER: 137:237717
 TITLE: Inhalant compositions containing anticholinergics and
 NK1 receptor antagonists
 INVENTOR(S): Meade, Christopher John Montague; Pairet, Michel;
 Pieper, Michael Paul
 PATENT ASSIGNEE(S): Boehringer Ingelheim Pharma K.-G., Germany
 SOURCE: PCT Int. Appl., 42 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 19
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002069944	A2	20020912	WO 2002-EP1987	20020226

WO 2002069944 A3 20031002

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW
 RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

DE 10111058 A1 20020912 DE 2001-10111058 20010308
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 EP 1370293 A2 20031217 EP 2002-719915 20020226
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
 JP 2004519484 T 20040702 JP 2002-569121 20020226
 MX 2003PA08051 A 20031204 MX 2003-PA8051 20030905
 AU 2006202723 A1 20060713 AU 2006-202723 20060626
 AU 2006202723 B2 20080626
 AU 2008202554 A1 20080703 AU 2008-202554 20080610

PRIORITY APPLN. INFO.:

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 EP 2001-200657 A 20010223
 AU 2002-308306 A3 20020222
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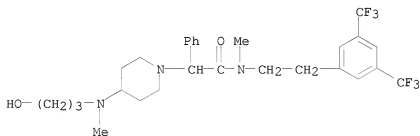
OTHER SOURCE(S): MARPAT 137:237717

IT 415916-92-6P 415917-07-6P 457910-79-1P
 457911-01-2P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (inhalant compns. contg. anticholinergics and NK1 receptor antagonists)

RN 415916-92-6 CAPLUS

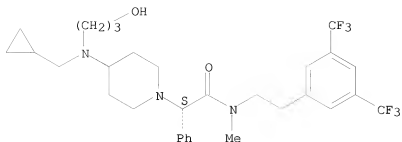
CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(3-hydroxypropyl)methylamino]-N-methyl-.alpha.-phenyl- (CA INDEX NAME)



RN 415917-07-6 CAPLUS

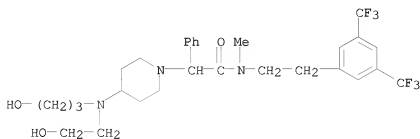
CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(cyclopropylmethyl)(3-hydroxypropyl)amino]-N-methyl-.alpha.-phenyl-, (.alpha.S)- (CA INDEX NAME)

Absolute stereochemistry.



RN 457910-79-1 CAPLUS

CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(2-hydroxyethyl)(3-hydroxypropyl)amino]-N-methyl-.alpha.-phenyl- (CA INDEX NAME)



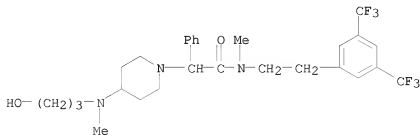
RN 457911-01-2 CAPLUS

CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(3-hydroxypropyl)methylamino]-N-methyl-.alpha.-phenyl-, (2E)-2-butenedioate (2:1) (salt) (9CI) (CA INDEX NAME)

CM 1

CRN 415916-92-6

CMF C28 H35 F6 N3 O2



CM 2

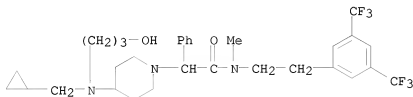
CRN 110-17-8

CMF C4 H4 O4

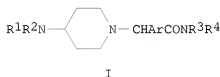
Double bond geometry as shown.



IT 457910-81-5
 RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (inhalant compns. contg. anticholinergics and NK1 receptor antagonists)
 RN 457910-81-5 CAPLUS
 CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-
 [(cyclopropylmethyl) (3-hydroxypropyl) amino]-N-methyl-.alpha.-phenyl- (CA
 INDEX NAME)



L4 ANSWER 6 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN
 TI 4-Aminopiperidinylacetamides as neurokinin antagonists
 GI



AB Title compds. I [R1 = (CH2)3OH, CH2CH(OH)CH2OH, cycloalkylmethyl; R2 = H, alkyl, hydroxyalkyl, CH2CH(OH)CH2OH, cycloalkylmethyl; R3 = (un)substituted Ph; R4 = H, alkyl, cycloalkyl, CH2CO2H, CH2CONH2. OH, phenylalkyl; Ar = (un)substituted Ph] were prepd. Thus, 1-benzyl-4-piperidinone was treated with H2N(CH2)3OH, N-methylated, debenzylated, and treated with 3,5-(F3C)2C6H3CH2CH2NMeCOCHPhO3SMe to give I [R1 = (CH2)3OH, R2 = R3 = Me, R4 = 3,5-(F3C)2C6H3CH2CH2]. At 0.2 .mu.Mol/kg iv in guinea pigs this compd. was effective in lowering blood pressure for > 360 min.

ACCESSION NUMBER: 2002:314907 CAPLUS
 DOCUMENT NUMBER: 136:340590
 TITLE: 4-Aminopiperidinylacetamides as neurokinin antagonists
 INVENTOR(S): Dollinger, Horst; Esser, Franz; Jung, Birgit; Schromm, Kurt; Speck, Georg
 PATENT ASSIGNEE(S): Boehringer Ingelheim Pharma K.-G., Germany
 SOURCE: PCT Int. Appl., 36 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002032865	A1	20020425	WO 2001-EP11906	20011016

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW

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US 6747044	B2	20040608		
CA 2426221	A1	20030417	CA 2001-2426221	20011016
EP 1328516	A1	20030723	EP 2001-987744	20011016

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JP 2004514658	T	20040520	JP 2002-536049	20011016
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PRIORITY APPLN. INFO.: DE 2000-10051320 A 20001017
US 2000-250541P P 20001201
WO 2001-EP11906 W 20011016

OTHER SOURCE(S): MARPAT 136:340590

IT 415917-04-3P 415917-07-6P 502422-75-5P

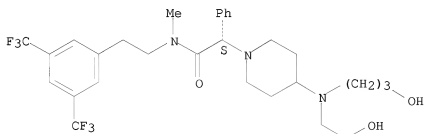
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of 4-aminopiperidinylacetamides as neurokinin antagonists)

RN 415917-04-3 CAPLUS

CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(2-hydroxyethyl)(3-hydroxypropyl)amino]-N-methyl-.alpha.-phenyl-, (.alpha.S)- (CA INDEX NAME)

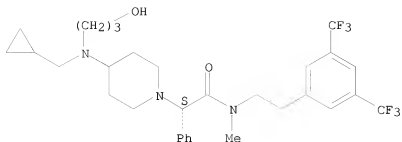
Absolute stereochemistry. Rotation (+).



RN 415917-07-6 CAPLUS

CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(cyclopropylmethyl)(3-hydroxypropyl)amino]-N-methyl-.alpha.-phenyl-, (.alpha.S)- (CA INDEX NAME)

Absolute stereochemistry.

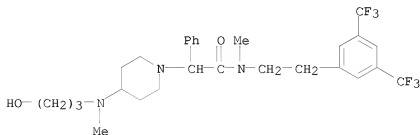


RN 502422-75-5 CAPLUS
 CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(3-hydroxypropyl)methylamino]-N-methyl-.alpha.-phenyl-, (2E)-2-butenedioate (2:3) (salt) (9CI) (CA INDEX NAME)

CM 1

CRN 415916-92-6

CMF C28 H35 F6 N3 O2

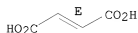


CM 2

CRN 110-17-8

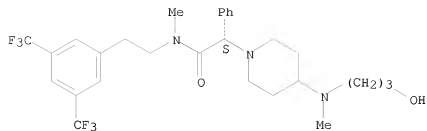
CMF C4 H4 O4

Double bond geometry as shown.



IT 415917-12-3P
 RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prepn. of 4-aminopiperidinylacetamides as neurokinin antagonists)
 RN 415917-12-3 CAPLUS
 CN 1-Piperidineacetamide, N-[2-[3,5-bis(trifluoromethyl)phenyl]ethyl]-4-[(3-hydroxypropyl)methylamino]-N-methyl-.alpha.-phenyl-, (.alpha.S)- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT:

2

THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT